Food and Drug Administration, HHS

- (ii) Function: Reviews proposed regulations for good manufacturing practices governing the methods used in, and the facilities and controls used for, the manufacture, packing, storage, and installation of devices, and makes recommendations on the feasibility and reasonableness of the proposed regulations.
- (3) Technical Electronic Product Radiation Safety Standards Committee.
 - (i) Date established: October 18, 1968.
- (ii) Function: Advises on technical feasibility, reasonableness, and practicability of performance standards for electronic products to control the emission of radiation under 42 U.S.C. 263f(f)(1)(A).
- (4) National Mammography Quality Assurance Advisory Committee.
 - (i) Date established: July 6, 1993.
- (ii) Function: Advises on developing appropriate quality standards and regulations for the use of mammography facilities.
- (e) National Center for Toxicological Research—Science Advisory Board.
 - (1) Date established: June 2, 1973
- (2) Function: Advises on establishment and implementation of a research program that will assist the Commissioner of Food and Drugs tofulfill regulatory responsibilities.
- (f) Center for Veterinary Medicine.
 Veterinary Medicine Advisory Committee.
 - (1) Date established: April 24, 1984.
- (2) Function: Reviews and evaluates available data concerning safety and effectiveness of marketed and investigational new animal drugs, feeds, and devices for use in the treatment and prevention of animal disease and increased animal production.
- (g) Center for Food Safety and Applied Nutrition—Food Advisory Committee.
- (1) Date established: December 15, 1991.
- (2) Function: The committee provides advice on emerging food safety, food science, and nutrition issues that FDA considers of primary importance in the next decade.

[54 FR 9036, Mar. 3, 1989]

EDITORIAL NOTE: For FEDERAL REGISTER citations affecting §14.100, see the List of CFR Sections Affected, which appears in the Finding Aids section of the printed volume and at www.fdsus.gov.

Subpart G—Technical Electronic Products Radiation Safety Standards Committee

§14.120 Establishment of the Technical Electronic Product Radiation Safety Standards Committee (TEPRSSC).

The Technical Electronic Product Radiation Safety Standards Committee (TEPRSSC), consisting of 15 members, is established in accordance with the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360kk(f)(1)(A)) to provide consultation before the Commissioner prescribes any performance standard for an electronic product.

[44 FR 22351, Apr. 13, 1979, as amended at 78 FR 17087, Mar. 20, 2013]

§ 14.122 Functions of TEPRSSC.

- (a) In performing its function of advising the Commissioner, TEPRSSC—
- (1) May propose electronic product radiation safety standards to the Commissioner for consideration;
- (2) Provides consultation to the Commissioner on all performance standards proposed for consideration under 21 U.S.C. 360kk; and
- (3) May make recommendations to the Commissioner on any other matters it deems necessary or appropriate in fulfilling the purposes of the act.
- (b) Responsibility for action on performance standards under 21 U.S.C. 360kk rests with the Commissioner, after receiving the advice of TEPRSSC.

 $[44\ {\rm FR}\ 22351,\ {\rm Apr.}\ 13,\ 1979,\ {\rm as}\ {\rm amended}\ {\rm at}\ 78\ {\rm FR}\ 17087,\ {\rm Mar.}\ 20,\ 2013]$

§ 14.125 Procedures of TEPRSSC.

- (a) When the Commissioner is considering promulgation of a performance standard for an electronic product, or an amendment of an existing standard, before issuing a proposed regulation in the FEDERAL REGISTER the Commissioner will submit to TEPRSSC the proposed standard or amendment under consideration, together with other relevant information to aid TEPRSSC in its deliberations.
- (b) The agenda and other material to be considered at any meeting will be sent to members whenever possible at least 2 weeks before the meeting.